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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/931,862	08/20/2001	Hae-Kyoung Kim	249/274	3541
7590 02/23/2004		•	EXAMINER	
LEE & STERBA, P.C. 1101 Wilson Boulevard			DOVE, TRACY MAE	
Suite 2000	Julevalu		ART UNIT	PAPER NUMBER
Arlington, VA 22209			1745	
			DATE MAILED 02/22/200	

DATE MAILED: 02/23/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

*		
	Application No.	Applicant(s)
	09/931,862	KIM, HAE-KYOUNG
Office Action Summary	Examiner	Art Unit
	Tracy Dove	1745
The MAILING DATE of this communication ap Period for Reply	ppears on the cover sheet with the	correspondence address
A SHORTENED STATUTORY PERIOD FOR REPL	VIS SET TO EXPIRE 2 MONTH	(S) EDOM
THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reg. If NO period for reply is specified above, the maximum statutory period. - Failure to reply within the set or extended period for reply will, by statuly any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	136(a). In no event, however, may a reply be tictly within the statutory minimum of thirty (30) date will apply and will expire SIX (6) MONTHS from the cause the application to become ABANDONI	mely filed ys will be considered timely. In the mailing date of this communication. ED (35 U.S.C. § 133).
Status		
1) Responsive to communication(s) filed on 17 L	December 2003.	
	s action is non-final.	
3) Since this application is in condition for allowa	ance except for formal matters, pr	osecution as to the merits is
closed in accordance with the practice under	Ex parte Quayle, 1935 C.D. 11, 4	53 O.G. 213.
Disposition of Claims		
4) Claim(s) 1-23 is/are pending in the application	٦.	
4a) Of the above claim(s) is/are withdra	awn from consideration.	
5) Claim(s) is/are allowed.		
6)⊠ Claim(s) <u>1-23</u> is/are rejected.		•
7) Claim(s) is/are objected to.		
8) Claim(s) are subject to restriction and/	or election requirement.	
Application Papers		
9)⊠ The specification is objected to by the Examin	er.	
10)⊠ The drawing(s) filed on 20 August 2001 is/are:	: a)⊠ accepted or b)⊡ objected	to by the Examiner.
Applicant may not request that any objection to the	e drawing(s) be held in abeyance. Se	e 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correct	ction is required if the drawing(s) is ob	ejected to. See 37 CFR 1.121(d).
11) The oath or declaration is objected to by the E	xaminer. Note the attached Office	e Action or form PTO-152.
Priority under 35 U.S.C. § 119		
 12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document)-(d) or (f).
2. Certified copies of the priority documen	ts have been received in Applicat	ion N o
3. Copies of the certified copies of the price	ority documents have been receive	ed in this National Stage
application from the International Burea	au (PCT Rule 17.2(a)).	
* See the attached detailed Office action for a list	t of the certified copies not receive	ed.
Attachment(s)		
D	4)	
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) 5) Notice of Informal F	Patent Application (PTO-152)
Paper No(s)/Mail Date	6) 🔛 Other:	

DETAILED ACTION

This Office Action is in response to the communication filed on 12/17/03. Applicant's argument have been considered, but are not persuasive. Claims 1-23 are pending are remain rejected in view of the prior art. This Action is made FINAL, as necessitated by amendment.

Specification

The amendment filed 12/17/03 is objected to under 35 U.S.C. 132 because it introduces new matter into the disclosure. 35 U.S.C. 132 states that no amendment shall introduce new matter into the disclosure of the invention. The added material which is not supported by the original disclosure is as follows: the specification as amended recites "a solid polymer electrolyte (SPE) fuel cell operable at *a temperature of room temperature to 200°C*", which does not appear to be supported by the specification as filed (or literature cited by the Examiner).

Applicant is required to cancel the new matter in the reply to this Office Action.

Claim Rejections - 35 USC § 112

The 35 U.S.C. 112, 2nd paragraph, rejections have been withdrawn.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1-3, 6-12 and 15-23 are rejected under 35 U.S.C. 102(b) as being anticipated by Bahar et al., US 5,635,041.

Regarding claims 1-3, Bahar teaches a composite membrane comprising a base material 4 and an ion exchange material/resin 2. The base material is a porous microstructure (porous support) and the ion exchange resin impregnates the membrane, i.e. base material (col. 3, lines 29-40). The ion exchange material may be comprised of at least in part a powder, such as but not limited to, carbon black, graphite, nickel, silica, titanium dioxide and platinum black (col. 2, lines 58-61). Optionally, the ion exchange materials may be complemented by finely divided powders or other (non-ionic) polymers to provide final composites. Such a finely divided powder may be selected from organic or inorganic compounds such as, but not limited to, carbon black, graphite, nickel, silica (SiO_2), titanium dioxide (TiO_2) or platinum black (catalyst). The powders provide specific added effects such as electrical conductivity, thermal conductivity, catalytic effects and/or enhanced or reduced reactant transport properties (col. 4, line 66-col. 5, line 8). Note silica and titanium dioxide are moisture retentive materials and platinum is a catalyst, as described in the instant specification (page 9, lines 9-21).

Regarding claims 6-7, a preferred base material is expanded polytetrafluoroethylene (ePTFE) having a porosity of greater than 35%, preferably between 70-95% (col. 3, lines 62-67).

Regarding claim 8, suitable ion exchange materials include perfluorinated sulfonic acid resin, perfluorinated carboxylic acid resin, polyvinyl alcohol, divinyl benzene, styrene-based polymers and metal salts with or without a polymer (col. 4, lines 58-63).

Regarding claims 9 and 23, a solution is prepared containing an ion exchange material (and optionally a finely divided powder). The solution may be applied to the base material by any conventional coating technique including roll coating, gravure coating, doctor coating, kiss

coating, dipping, brushing, painting or spraying so long as the liquid solution is able to penetrate the interstices and interior volume of the base material (col. 6, lines 19-27).

Regarding claims 10-12 and 15-18, the composite membrane may be used in a fuel cell (claim 4). Ion exchange membranes are used in polymer electrolyte fuel cells as solid electrolytes (col. 1, lines 14-15). The composite membrane of Bahar may be used in various applications, including fuel cells and batteries (col. 3, lines 41-44).

Regarding claims 19-22, a direct methanol fuel cell (DMFC) has the same structure as the polymer electrolyte membrane fuel cell (PEMFC), but uses liquid methanol, instead of hydrogen, as a fuel source (see page 3, lines 13-14 of the instant specification "Description of Related Art). Thus, the direct methanol fuel cell of claim 19 is a polymer electrolyte fuel cell (taught by Bahar). Note that whether methanol or hydrogen is used as the fuel source, the fuel cell is a polymer electrolyte type fuel cell (the terminology of the preamble does not limit the claimed structure MPEP 2111.02).

Thus, the claims are anticipated.

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Claims 1, 2, 5-11 and 14-23 are rejected under 35 U.S.C. 102(b) as being anticipated by Grot et al., US 5,919,583.

Regarding claims 1, 2, 6, 7, 10, 11, 15, 16, 19 and 21, Grot teaches a cation exchange membrane made from a polymer having cation exchange groups and containing inorganic filler. The membrane exhibits reduced fuel crossover for fuel cells employing direct feed organic fuels such as methanol (see abstract). Preferably the inorganic filler is an inorganic proton conductor selected from the group consisting of titanium dioxide, tin and hydrogen mordenite, oxides and

phosphates of zirconium, and mixtures thereof. The inorganic proton conductor comprises 2-30 wt% of the membrane (col. 2, lines 25-38). The membrane may optionally include a porous support for improving mechanical properties. The porous support may be a polyolefin (polyethylene or polypropylene) or polytetrafluoroethylene (PTFE) having at least 40% porosity (col. 5, lines 1-31). A membrane can be made using a porous support by coating cation exchange polymer on the support so that the coating is on the outside surfaces as well as being distributed through the internal pores (impregnates) of the support (col. 5, lines 32-33). The inorganic filler is dispersed in the membrane (impregnates) and may further be a zeolite material (col. 5, lines 58-63). Note titanium dioxide, zirconium oxide, mordenite and zeolite are moisture retentive materials, as described in the instant specification (page 9, lines 9-21).

Regarding claims 5, 8, 14, 17, 20 and 22, the cation exchange groups of the polymer are selected from the group consisting of sulfonate, carboxylate, phosphonate, imide, sulfonimide and sulfonamide. In a preferred embodiment, highly fluorinated polymer with sulfonate groups is employed (col. 2, lines 39-50). The term "sulfonate groups" is intended to refer either to sulfonic acid groups or alkali metal or ammonium salts of sulfonic acid groups (col. 3, lines 57-60). Example 2 teaches a solution containing a sulfonated perfluorocarbon copolymer having as a perfluorocarbon backbone and side chains -O-CF₂CF(CF₃)-O-CF₂CF₂SO₃H in hydrogen ion form and which has an equivalent weight of about 1080. Tin mordenite is added to the solution and the solution is poured onto a polytetrafluoroethylene sheet substrate (porous support).

Regarding claims 9, 18 and 23, a solution of an inorganic filler and a polymer ionic form can be used to apply a coating to a porous support to form the membrane (col. 6, lines 42-46). The polymer is distributed through the internal pores of the support (col. 5, lines 32-35) and the

Application/Control Number: 09/931,862

Art Unit: 1745

inorganic filler is incorporated into the membrane (col. 6, line 40). Thus, the membrane of Grot is formed by impregnating the porous support with a composition of ion-exchange polymer and the inorganic filler (reinforcing agent).

Thus the claims are anticipated.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 1-4, 8, 10-13, 17 and 19 are rejected under 35 U.S.C. 103(a) as being unpatentable over Watanabe et al., US 5,766,787 in view of Grot et al., US 5,919,583.

Watanbe teaches a solid polymer electrolyte fuel cell comprising a solid polymer electrolyte membrane incorporating 5.8 wt% platinum catalyst and 5 wt% silica in Nafion (perfluorocarbon sulfonic acid cation exchange resin) or 5.8 wt% platinum catalyst and 5 wt% titania (TiO₂) in Nafion. See col. 6, lines 40-48 and col. 8, lines 23-64. Thus, the platinum catalyst is about 54 wt% and the silica (or titania) is about 46 wt% of the total amount of catalyst plus metal oxide (reinforcing agent) contained in the polymer electrolyte membrane. The membrane comprises 0.01-80 wt% of at least one metal catalyst (Pt, Au, Pd, Rh, Ir and/or Ru) and 0.01-50wt% of particles and/or fibers of at least one metal oxide (silica, titania and/or zirconia). See col. 3, lines 29-42. Methanol gas and oxygen gas may be used as the reactant gases for the fuel cell (col. 3, lines 57-59).

Watanabe does not explicitly state the polymer electrolyte membrane includes a porous support.

However, Grot teaches a cation exchange membrane made from a polymer having cation exchange groups and containing inorganic filler. The membrane exhibits reduced fuel crossover for fuel cells employing direct feed organic fuels such as methanol (see abstract). Preferably the inorganic filler is an inorganic proton conductor selected from the group consisting of titanium dioxide, tin and hydrogen mordenite, oxides and phosphates of zirconium, and mixtures thereof. The inorganic proton conductor comprises 2-30 wt% of the membrane (col. 2, lines 25-38). The membrane may optionally include a porous support for improving mechanical properties, for decreasing cost and/or other reasons. The porous support may be a polyolefin (polyethylene or polypropylene) or polytetrafluoroethylene (PTFE) having at least 40% porosity (col. 5, lines 1-31). A membrane can be made using a porous support by coating cation exchange polymer on the support so that the coating is on the outside surfaces as well as being distributed through the internal pores (impregnates) of the support (col. 5, lines 32-33). The inorganic filler is dispersed in the membrane (impregnates) and may further be a zeolite material (col. 5, lines 58-63). Note titanium dioxide, zirconium oxide, mordenite and zeolite are moisture retentive materials, as described in the instant specification (page 9, lines 9-21).

Therefore, the invention as a whole would have been obvious to one having ordinary skill in the art at the time the invention was made because one of skill would have been motivated to incorporate a porous support in the polymer electrolyte membrane of Watanabe in order to improve the mechanical properties and/or decrease the cost of the membrane (see Grot col. 5, lines 1-3). Grot teaches that the polymer electrolyte membranes optionally include a porous

support. Therefore, one of skill in the art would be motivated to provide a porous support in the polymer electrolyte membrane of Watanabe in order to improve the mechanical properties of the membrane and/or to decrease the cost of the membrane. Grot teaches membranes containing a cation exchange polymer and a reinforcing agent (as defined by the instant specification) may or may not include a porous support. Both Watanabe and Grot teach direct methanol fuel cells.

Response to Arguments

Applicant's arguments filed 12/17/03 have been fully considered but they are not persuasive.

Specification

Applicant states that the operating temperature range for a polymer electrolyte fuel cell (PEFC) has been amended to recite a range of room temperature to 200°C as disclosed on table 4-1 of Fuel Cells and their Applications (1996), which was cited by the Examiner. However, table 4-1 discloses a temperature range of 50-80°C for a PEFC.

35 U.S.C. 102(b) in view of Bahar

Applicant argues Bahar fails to disclose a reinforced composite ionic conductive polymer membrane as recited in the present claims. Specifically, Applicant argues Bahar fails to disclose "a separate reinforcing agent". Examiner points out that the claims do not require "a separate reinforcing agent", as asserted by Applicant. At least claims 9, 18 and 23 teach the reinforcing agent is mixed with the ion-exchange polymer because the claims recite "a composition of the ion-exchange polymer and the reinforcing agent" (claims 9 and 18) and "a mixture of an ionexchange polymer and a reinforcing agent" (claim 23). Thus, Applicant's arguments are not persuasive.

35 U.S.C. 102(b) in view of Grot

Applicant argues, like Bahar, Grot fails to disclose the use of a reinforcing agent that is separate from the ion-exchange polymer. As indicated above, the claims do not require "a separate reinforcing agent", as asserted by Applicant. At least claims 9, 18 and 23 teach the reinforcing agent is mixed with the ion-exchange polymer because the claims recite "a composition of the ion-exchange polymer and the reinforcing agent" (claims 9 and 18) and "a mixture of an ion-exchange polymer and a reinforcing agent" (claim 23). Thus, Applicant's arguments are not persuasive.

35 U.S.C. 103(a) over Watanabe in view of Grot

Applicant argues neither Watanabe nor Grot teaches adding a separate reinforcing agent that impregnates the porous support as recited in the present claims. However, the claims do not require "a separate reinforcing agent", as asserted by Applicant. At least claims 9, 18 and 23 teach the reinforcing agent is mixed with the ion-exchange polymer because the claims recite "a composition of the ion-exchange polymer and the reinforcing agent" (claims 9 and 18) and "a mixture of an ion-exchange polymer and a reinforcing agent" (claim 23). Thus, Applicant's arguments are not persuasive.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO

MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Tracy Dove whose telephone number is 571-272-1285. The examiner can normally be reached on Monday-Thursday (9:00-7:30).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Pat Ryan can be reached on 571-272-1292. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

February 12, 2004